

Measurements of slow tissue dynamics with short-separation speckle contrast optical spectroscopy: supplement

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1. CHOOSING THE g_1 MODELS FOR SLOW TISSUE DYNAMICS

The scattering regimes and particle motion types for blood flow dynamics have been well discussed in the literature[1, 2] and can be adapted to characterize the slow tissue dynamics we are investigating. The field autocorrelation function $g_1 = \exp[-(\frac{\tau}{\tau_{c2}})^n]$ with $n = 0.5, 1, 2$ was applied to derive the corresponding speckle contrast models [2]. We evaluated the post-euthanasia mouse data at $T_{exp} > 1s$ in order to reduce the effects of blood flow. Then we fitted the data with different speckle contrast models with a single decorrelation time τ_{c2} as described in [2]. Fig. S1 shows that the speckle contrast model based on $g_1 = \exp[-(\frac{\tau}{\tau_{c2}})^n]$ with $n = 1$ has the best fitting accuracy compared to models with $n = 0.5$ and $n = 2$, suggesting that the slow tissue dynamics measured with ss-SCOS is dominated by single scattering with unordered motion or multiple scattering with ordered motion [1]. Note that the ROI in the brain surface was selected in the region of parenchyma with small vessels.

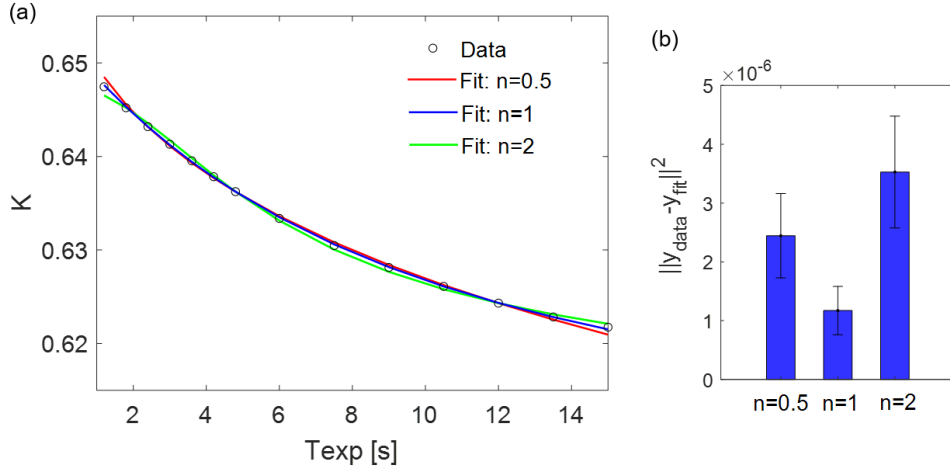


Fig. S1. Comparison of slow tissue dynamics fitted with different g_1 models. (a) Post-euthanasia mouse data (50 minutes after pentobarbital overdose injection) and the fitting curves with different g_1 models. (b) Fitting residual norm $\|y_{data} - y_{fit}\|^2$ for different g_1 models. The error bar represents the standard error of 5 repetitive measurements.

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